

FORM PTO-1390
(REV. 11-2000)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NUMBER

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)
CONCERNING A FILING UNDER 35 U.S.C. 371

264/264

U.S. APPLICATION NO. (If known, see 37 CFR 1.5)

09/868289

INTERNATIONAL APPLICATION NO.

PCT/EP99/10040

INTERNATIONAL FILING DATE

December 17, 1999

PRIORITY DATE CLAIMED

December 17, 1998

TITLE OF INVENTION

THERMOSETTING POLY URETHANE/UREA-FORMING COMPOSITIONS

APPLICANT(S) FOR DO/EO/US

Didier ARNOUX; Crispin Frank Maxwell JONES; Ronald Owen ROSENBERG

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:

1. ☒ This is a **FIRST** submission of items concerning a filing under 35 U.S.C. 371.
2. ☐ This is a **SECOND** or **SUBSEQUENT** submission of items concerning a filing under 35 U.S.C. 371.
3. ☐ This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.
4. ☒ The US has been elected by the expiration of 19 months from the priority date (Article 31).
5. ☒ A copy of the International Application as filed (35 U.S.C. 371(c)(2))
 - a. ☐ is attached hereto (required only if not communicated by the International Bureau)
 - b. ☒ has been communicated by the International Bureau
 - c. ☐ is not required, as the application was filed in the United States Receiving Office (RO/US).
6. ☐ An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).
 - a. ☐ is attached hereto.
 - b. ☐ has been previously submitted under 35 U.S.C. 154(d)(4)
7. ☒ Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))
 - a. ☐ are attached hereto (required only if not communicated by the International Bureau).
 - b. ☐ have been communicated by the International Bureau.
 - c. ☐ have not been made; however, the time limit for making such amendments has NOT expired.
 - d. ☒ have not been made and will not be made.
8. ☐ An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).
9. ☐ An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).
10. ☐ An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).

Items 11 to 20 below concern document(s) or information included:

11. ☐ An Information Disclosure Statement under 37 CFR 1.97 and 1.98.
12. ☐ An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.
13. ☐ A FIRST preliminary amendment.
14. ☐ A SECOND or SUBSEQUENT preliminary amendment.
15. ☐ A substitute specification.
16. ☐ A change of power of attorney and/or address letter.
17. ☐ A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.
18. ☒ A second copy of the published international application under 35 U.S.C. 154(d)(4).
19. ☐ A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).
20. ☒ Other items or information: **Form PCT/IB/308**

15 JUN 2001

U.S. APPLICATION NO. (if known, see 37 CFR 1.53) 09/868289		INTERNATIONAL APPLICATION NO. PCT/EP99/10040	ATTORNEY'S DOCKET NUMBER 264/264	
21. <input checked="" type="checkbox"/> The following fees are submitted: BASIC NATIONAL FEE (37 CFR 1.492(a)(1) – (5)): Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$1000.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$860.00 International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$710.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$690.00 International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00 ENTER APPROPRIATE BASIC FEE AMOUNT =			CALCULATIONS PTO USE ONLY	
Surcharge of \$130.00 for furnishing the oath or declaration later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).			\$	
CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$
Total claims	19 - 20 =	0	x \$18.00	\$
Independent claims	3 - 3 =	0	x \$80.00	\$
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$270.00	\$
TOTAL OF ABOVE CALCULATIONS =			\$860.00	
<input type="checkbox"/> Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.			\$	
SUBTOTAL =			\$860.00	
Processing fee of \$130.00 for furnishing the English translation later than <input checked="" type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).			\$	
TOTAL NATIONAL FEE =			\$860.00	
Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property			\$	
TOTAL FEES ENCLOSED =			\$	
			Amount to be refunded:	\$
			charged:	\$860.00
a. <input type="checkbox"/> A check in the amount of \$ to cover the above fees is enclosed.				
b. <input checked="" type="checkbox"/> Please charge my Deposit Account No. 12-2475 in the amount of \$860.00 to cover the above fees. A duplicate copy of this sheet is enclosed.				
c. <input checked="" type="checkbox"/> The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 12-2475. A duplicate copy of this sheet is enclosed.				
d. <input type="checkbox"/> Fees are to be charged to a credit card. WARNING: Information on this form may become public. Credit card information should not be included on this form. Provide credit card information and authorization on PTO-2038.				
SEND ALL CORRESPONDENCE TO: LYON & LYON 633 WEST FIFTH STREET, SUITE 4700 LOS ANGELES, CALIFORNIA 90071-2066 — (213) 489-1600 KRISTIN H. NEUMAN DATE: <u>June 15, 2001</u>			<u>Kristin H. Neuman</u> SIGNATURE NAME <u>KRISTIN H. NEUMAN</u> REGISTRATION NUMBER <u>35,530</u>	

THERMOSETTING POLY URETHANE/UREA-FORMING COMPOSITIONS

This invention relates to casting polyurethane and/or poly urethane/urea-forming products capable of room temperature curing. More particularly, the invention is drawn to liquid compositions containing isocyanate-functional prepolymers, an aromatic amine curative therefore and a non-volatile phosphate or phthalate plasticizer. The selected poly urethane/urea-forming compositions produce, after room-temperature cure, non-sweating, non-distorting castings or boards with superior elasticity, toughness, tear strength and abrasion resistance.

BACKGROUND OF THE INVENTION

Aromatic polyisocyanates are well known and are widely used in the preparation of polyurethane and poly urethane/urea elastomers. These aromatic diisocyanates generally include compositions such as 2,4-toluene diisocyanate, 2,6-toluene diisocyanate, 4,4'-methylene bis (phenylisocyanate) and the like. In the preparation of polyurethane and polyurethane/urea elastomers, the aromatic diisocyanates are reacted with a long chain (high molecular weight) polyol to produce a prepolymer containing free isocyanate groups which then may be chain extended with a short chain (low molecular weight) polyol or aromatic diamine to form a polyurethane or polyurethane/urea elastomer. Long chain, high molecular weight polyols, e.g. those having a molecular weight of above 250, are generally utilized to form the prepolymer and the chain extender is generally a short chain polyol, e.g., C₂-C₁₀ polyol, or an aromatic diamine. The long chain, high molecular weight polyol provides flexibility and elastomeric properties to the resin, while the short chain polyol or aromatic diamine provides chain extension or cross-links and adds toughness and rigidity to the resulting elastomeric polymer.

A major problem with mononuclear aromatic diisocyanates, such as, toluene diisocyanate, relates to the perceived health risks and their volatility. It is known that residual toluene diisocyanate (free toluene diisocyanate) in a prepolymer mixture can be reduced by lowering the isocyanate/hydroxyl ratio of the prepolymer mixture. This modification, however, has a detrimental effect on processing when the prepolymer is chain extended (or cured), namely, the hardness build up rate decreases very significantly, which leads to an

extended demolding time. The prepolymer viscosity also builds up to levels which prevent processability at ambient temperatures.

Alternatively, it is also known that residual toluene diisocyanate may be physically removed from a prepolymer by vacuum distillation or other methods.

It is also known that by incorporating 2,4 toluene diisocyanate dimer into a prepolymer mix, a low NCO/OH ratio can be used to obtain the desired low residual free toluene diisocyanate content while maintaining the desired hardness build up rate. Such a composition is described in U.S. Pat. No. 5,077,371, which is incorporated herein by reference. The prepolymer's viscosity is measured at 100°C and cured at temperatures in excess of 100°C using 4,4'-methylene-bis(3-chloro)aniline. No phosphate ester plasticizers or room temperature curing are described or contemplated by the teachings in the '371 Patent. There is a need for a poly urethane/urea forming composition having a low free toluene diisocyanate content that is capable of being cast and cured at room temperatures.

Plasticizers are commonly used for polyurethane foaming compositions to modify the foam properties or ease processing as described in U.S. Pat. No. 5,817,860. Examples of plasticizers set forth therein are dioctyl phthalate, diisooctyl phthalate, dimethyl phthalate, dibutyl phthalate, tributyl phosphate, triphenyl phosphate, cresyl diphenyl phosphate, halogenated biphenyls and aromatic oils.

U.S. Pat. No. 5,688,892 describes a waterproofing formulation based upon a cold setting polyurethane formulation using a selected aromatic diamine curative and plasticizers and a toluene diisocyanate reacted with a polyoxypropylene or polyoxyethylene polyol. The plasticizer used therein is dioctyl phthalate.

Published European patent application 829,497 describes a polyurethane casting composition based on an isocyanate or isocyanate-function prepolymer, selected aromatic amine curative with a relatively low polarity and hydrocarbon oils as a plasticizer.

Many conventional plasticizers, however, are either too volatile, which leads to dimensional instability, or become physically incompatible after cure, or have insufficient viscosity reducing effect. The present invention overcomes these obstacles to produce a desired

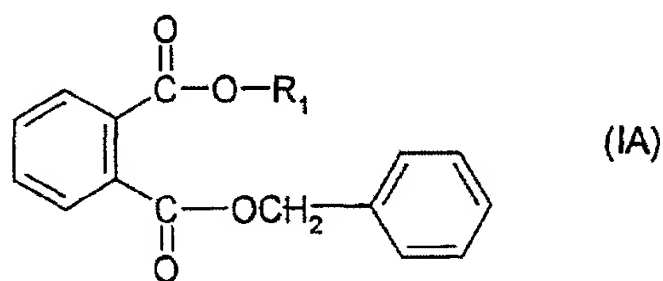
room temperature curable poly urethane/urea-forming product having a low free toluene diisocyanate content.

SUMMARY OF THE INVENTION

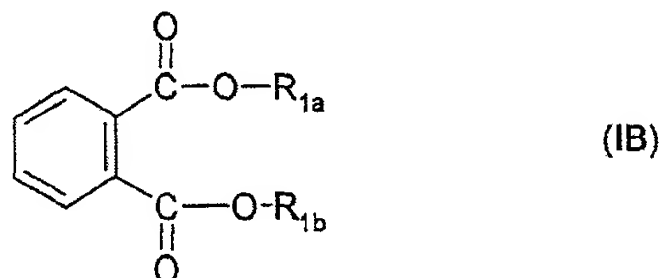
The present invention is, in one embodiment, drawn to a poly urethane/urea-forming casting composition that can be cast and cured at temperatures between 15 and 35°C using a reaction mixture of (a) an isocyanate component or an isocyanate functional prepolymer having at least two isocyanate groups per molecule that contains or has been reacted with polytetramethylene glycol, (b) an aromatic amine curative, and (c) a phosphate ester or phthalate ester having a vapor pressure of less than 100 mPa at 25°C. Component (a) is preferably a low free toluene diisocyanate prepolymer blend. Component (a) more preferably is a prepolymer blend that is a reaction mixture of an organic diisocyanate and polytetramethylene glycol. The casting composition can further include a polyether- and/or polyester polyol having a number average molecular weight of at least 250.

The aromatic amine curative is preferably selected from the group consisting of diethyl toluene diamine, tertiary butyl toluene diamine, dimethylthiotoluene diamine, , and 1,2-bis(2-aminophenylthio)ethane.

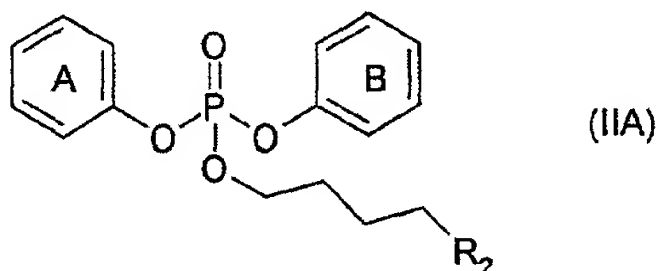
The phosphate ester or phthalate ester is preferably represented by formulae (IA), (IB), (IIA), (IIB) or (III):



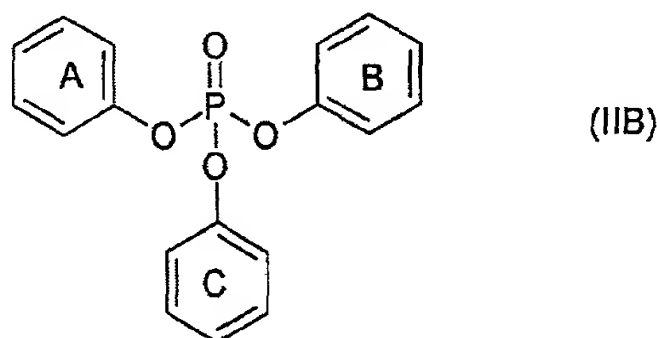
wherein R₁ is unsubstituted or alkyl-substituted C₃-C₁₂alkyl,



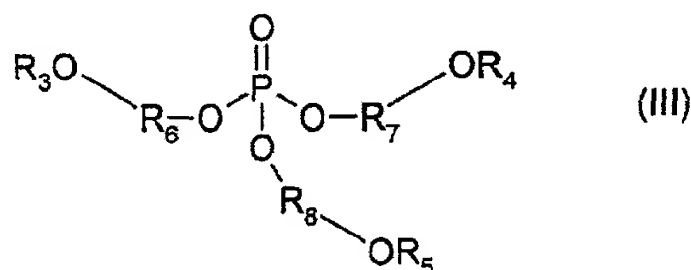
wherein R_{1a} and R_{1b} , independently of one another are unsubstituted or alkyl-substituted C_5 - C_{12} alkyl; or phosphate esters according to formulae (IIA) and (IIB)



wherein R_2 is hydrogen or unsubstituted or alkyl-substituted C_2 - C_6 alkyl, and aromatic rings A and B independently of one another can have one or more alkyl substitutions;



wherein aromatic rings A, B and C independently of one another can have one or more alkyl substitutions, or formula (III)

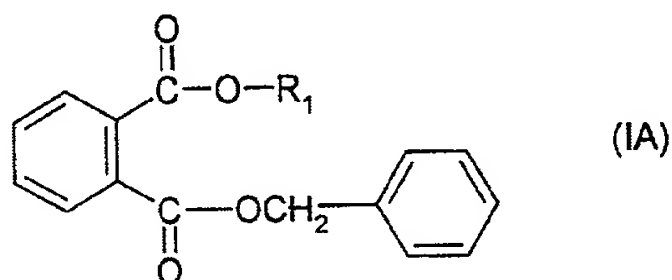


wherein R_3 , R_4 , R_5 , independently from one another, are unsubstituted or alkyl substituted C_1 - C_5 alkyl and R_6 , R_7 and R_8 are independently of one another are unsubstituted or alkyl-substituted C_1 - C_5 alkylene. More preferably, component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.

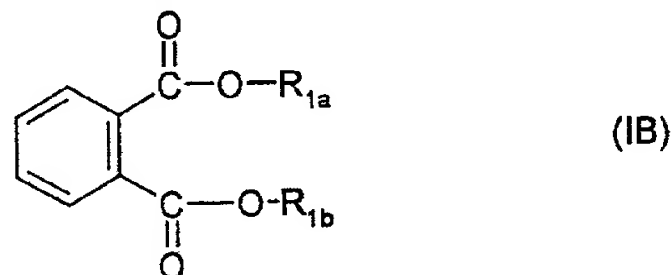
The present invention, in a further embodiment, is drawn to a cast polyurethane obtainable by a process comprising reacting (a) an isocyanate component or an isocyanate functional prepolymer having at least two isocyanate groups per molecule that contains or has been reacted with polytetramethylene glycol, (b) an aromatic amine curative, and (c) a phosphate ester or phthalate ester having a vapor pressure of less than 100 mPa at 25°C. Component (a) can be a prepolymer blend resulting from a reaction mixture of an organic diisocyanate

and polytetramethylene glycol. The aromatic amine curative used to obtain cast polyurethane is selected from the group consisting of diethyl toluene diamine, tertiary butyl toluene diamine, dimethylthiotoluene diamine, and 1,2-bis(2-aminophenylthio)ethane. The reactants for obtaining the desired cast polyurethane can further include a polyether- and/or polyester polyol having a number average molecular weight of at least 250.

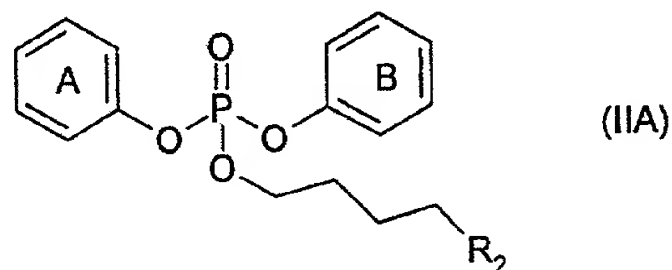
The phosphate ester or phthalate ester used for this process can preferably be represented by formulae (IA), (IB), (IIA), (IIB) or (III):



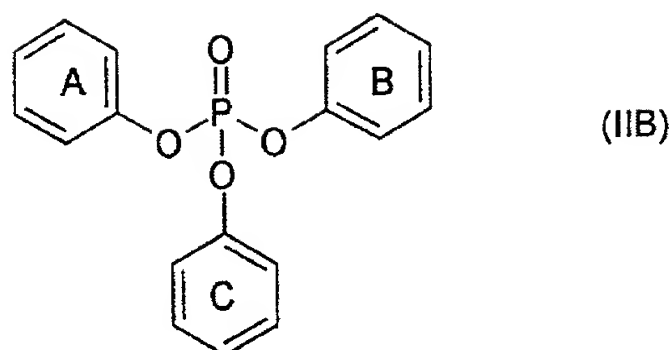
wherein R_1 is unsubstituted or alkyl-substituted C_3 - C_{12} alkyl,



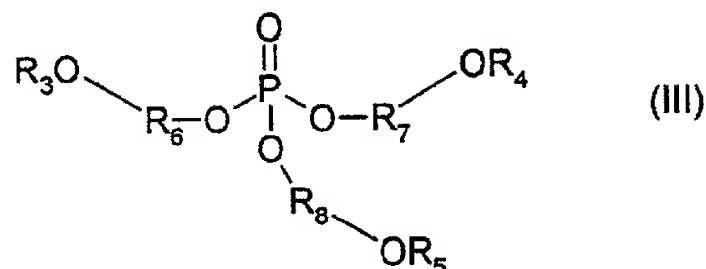
wherein R_{1a} and R_{1b} , independently of one another are unsubstituted or alkyl-substituted C_5 - C_{12} alkyl, or phosphate esters according to formulae (IIA) and (IIB)



wherein R_2 is hydrogen or unsubstituted or alkyl-substituted C_2 - C_6 alkyl, and aromatic rings A and B independently of one another can have one or more alkyl substitutions;



wherein aromatic rings A, B and C independently of one another can have one or more alkyl substitutions, or formula (III)



wherein R₃, R₄, R₅, independently from one another, are unsubstituted or alkyl substituted C₁-C₅ alkyl and R₆, R₇ and R₈ are independently of one another are unsubstituted or alkyl-substituted C₁-C₅alkylene. More preferably, component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.

The present invention is drawn to a still further embodiment of a process for curing a poly urethane/urea-forming composition comprising contacting said poly urethane/urea-forming composition at a temperature between 15 and 35°C with (a) an aromatic amine curative having at least two primary amine groups, and (b) a plasticizer having a vapor pressure of less than 100 mPa at 25°C with said poly urethane/urea-forming composition.

DETAILED DESCRIPTION OF THE INVENTION

The isocyanate used in this invention is an isocyanate having an average functionality of two or more isocyanate groups per molecule. Examples of suitable diisocyanates are 2,4-toluene diisocyanate, 4,4-diphenylmethane diisocyanate, mixtures of diphenylmethane diisocyanate, paraphenyldiisocyanate, isophoronediiisocyanate, 4,4'-methylene-bis(cyclohexylisocyanate), naphthalene diisocyanate and hexamethylene diisocyanate and chain extended prepolymer blends. An alkylene diamine or diol can be reacted with an excess of the isocyanate component to produce such a chain extended prepolymer blends.

In a preferred embodiment, a low-free toluene diisocyanate prepolymer is made by reacting toluene diisocyanate with high molecular weight aliphatic polyester or polyether polyols to produce a prepolymer having a toluene diisocyanate content below 0.4% by weight, optionally by removal of excess toluene diisocyanate. Optionally, a mixture of a low molecular weight polyol (number average molecular weight less than 250) and a high molecular weight polyol (number average molecular weight of at least 250) can also be reacted with the toluene diisocyanate.

Representative toluene diisocyanates include the two main isomers, 2,4- and 2,6-diisocyanate and optionally, a small amount of the ortho isomers, the 2,3- and 3,4-isomers. Commercially, toluene diisocyanate is found as a 65:35, 80:20 or 99:1 isomer mix of the 2,4- and 2,6-isomer by weight and optionally from 0-5% by weight of the ortho isomers. An isomer mix is preferred within a range of from about 65-100% of the 2,4-isomer and the balance (0-35%) being essentially the 2,6-isomer. The most preferred range of the 2,6-isomer is 20-35%.

The 2,4-toluene diisocyanate dimer can be added to the selected organic diisocyanate(s) in an amount ranging up to the dimer's solubility limit at about 80°C to form an isocyanate blend. Preferably the dimer is present at up to about 6.0% by weight of the isocyanate blend, that is, the mixture of the toluene diisocyanate dimer and the selected organic diisocyanate(s). More preferably up to about 4.0% by weight of dimer may be present in the isocyanate blend.

High molecular weight polyols, particularly polyether polyols or polyester polyols having a number average molecular weight of at least 250, can be used to prepare the prepolymer of the instant invention. Polyols having a number average molecular weight of about 650 to 3000 are preferred, with polyols having number average molecular weights of 1000 being the most preferred. The number average molecular weight of the high molecular weight polyol can be as high as 10,000 or as low as 250. Preferred polyether polyols are polyalkyleneether polyols represented by the general formula $\text{HO}(\text{RO})_n\text{H}$, wherein R is an alkylene radical and n is an integer large enough that the polyether polyol has a number average molecular weight of at least 250. These polyalkyleneether polyols are well-known components of polyurethane products and can be prepared by the polymerization of cyclic

ethers such as alkylene oxides and glycols, dihydroxyethers, and the like by known methods. A particularly preferred high molecular weight polyol is polytetramethylene glycol.

Polyester polyols are prepared by reaction of dibasic acids (usually adipic acid but other components such as sebacic or phthalic acid may be present) with diols such as ethylene glycol, 1,2-propylene glycol, 1,4-butylene glycol and diethylene glycol, etc., where linear polymer segments are required, or include units of higher functionality such as glycerol, trimethylol propane, pentaerythritol, sorbitol, etc., if chain branching or ultimate cross-linking is sought. Some polyester polyols also employ caprolactone and dimerized unsaturated fatty acids in their manufacture. Another type of polyester which is of interest is that obtained by the addition polymerization of ϵ -caprolactone in the presence of an initiator. Other polyols that can be used are those that have at least two hydroxyl groups and whose basic backbone is obtained by polymerization or copolymerization of such monomers as butadiene and isoprene monomers.

The initial polyol portion for making a prepolymer blend of the instant invention can be combination of high molecular weight polyol, as previously described, and a low molecular weight polyol. An aliphatic glycol is the preferred low molecular weight polyol. Suitable aliphatic polyols are ethylene glycol, diethylene glycol, dipropylene glycol, neopentyl glycol, 1,3-butanediol, 1,4-butanediol, and the like. The most preferred low molecular weight polyol is 1,4-butanediol. In general, the weight of the low molecular weight polyol should be no more than 20% of the combination of high molecular weight polyol and low molecular weight polyol. The preferred range is 0 to 15% of the combination; more preferred is 0-8%.

The prepolymers are preferably prepared by loading toluene diisocyanate, then adding the polyol or initial polyol blend, maintaining the temperature at from room temperature to temperatures as high as 150°C for sufficient time to react all of the available hydroxyl groups, then removing unreacted toluene diisocyanate by vacuum distillation or other physical means. Preferred reaction temperatures are 50°C to 100°C; more preferred temperatures are 50°C to 85°C. The product is poured into containers under a nitrogen flush and stored at room temperature. A particularly preferred prepolymer is commercially available from Uniroyal Corporation under the tradename Adiprene® LF 750D.

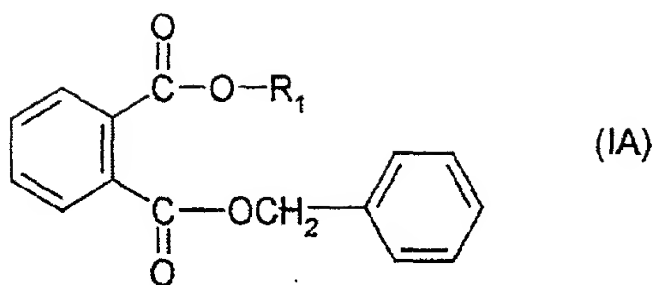
The stoichiometric ratio of isocyanato groups to hydroxyl groups in the reactants should preferably be from 2/1 to 30/1 although somewhat lower and higher ratios are permissible. When the ratio is much lower, the molecular weight of the isocyanato terminated polyurethane becomes so large (due to formation of oligomers with more than one polyol moiety) that the viscosity of the mass makes mixing of chain extenders into the prepolymer relatively more difficult. At the other extreme, an excess approaching the 30/1 ratio will result in high levels of free diisocyanate in the mixture (with higher removal costs). Therefore, the preferred range is 2.5/1 to 10/1.

The curative used with the prepolymer described above is an organic aromatic polyamine having two or more amine groups. The organic aromatic polyamine can be combined with polyether- and/or polyester-polyols described above to modify the final product or cure characteristics. Of course, known catalysts can be used in conjunction with the curative if necessary.

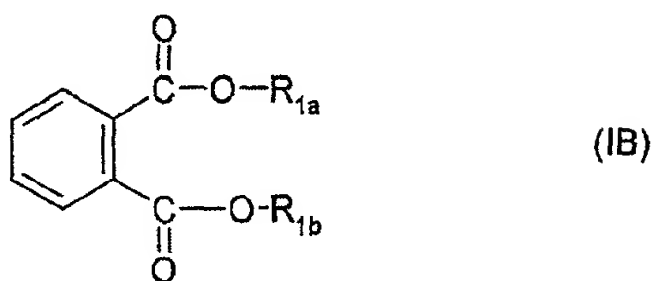
Examples of aromatic polyamine materials are: diethyl toluene diamine (DETDA), tertiary butyl toluene diamine (TBTDA), dimethylthiotoluene diamine (Ethacure™ 300) from Albermarle Corporation, trimethylene glycol di-p-aminobenzoate (Vibracure™ A157 from Uniroyal Chemical Company or Versalink™ 740 from AirProducts and Chemicals Inc.), and 1,2-bis(2-aminophenylthio)ethane (Cyanacure from American Cyanamid Company). The most preferred aromatic polyamine is dimethylthiotoluene diamine.

The castable composition combines the isocyanate component and/or prepolymer blend and curative described above with a plasticizing agent having a vapor pressure at 25°C of less than 100 mPa and/or an evaporation rate of <40% after 24 hours at 87°C according to ASTM 1203-67. The preferred plasticizing agents are phosphate or phthalate esters having a vapor pressure at 25°C of less than 100 mPa.

Particularly preferred plasticizing agents are alkyl benzyl phthalates monomers according to formulae (IA) and (IB)

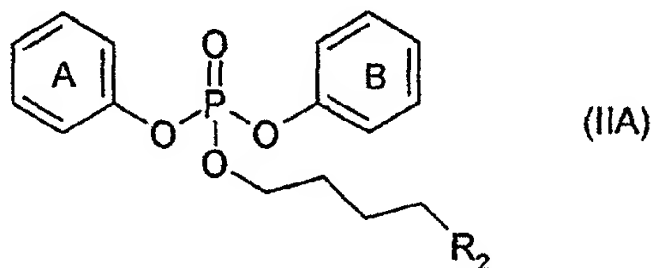


wherein R_1 is unsubstituted or alkyl-substituted C_3 - C_{12} alkyl,

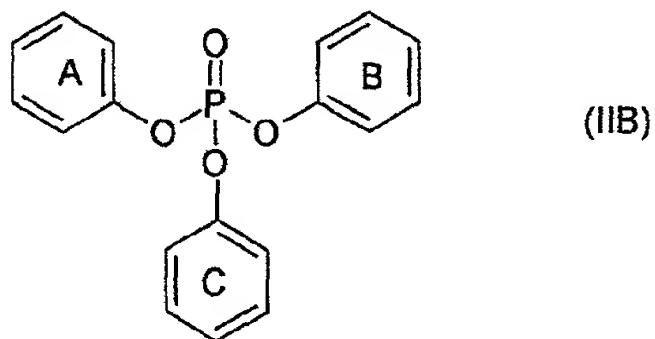


wherein R_{1a} and R_{1b} , independently of one another are unsubstituted or alkyl-substituted C_5 - C_{12} alkyl, preferably C_9 - C_{12} alkyl;

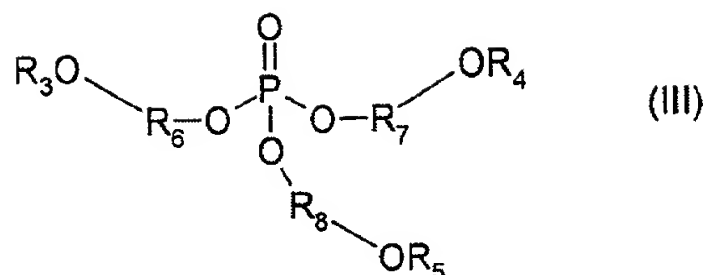
or phosphate esters according to formulae (IIA) and (IIB)



wherein R_2 is hydrogen or unsubstituted or alkyl-substituted C_2 - C_6 alkyl, and aromatic rings A and B independently of one another can have one or more alkyl substitutions;



wherein aromatic rings A, B and C independently of one another can have one or more alkyl substitutions,
or formula (III)



wherein R_3 , R_4 , R_5 , independently from one another, are unsubstituted or alkyl substituted C_1 - C_5 alkyl and R_6 , R_7 and R_8 are independently of one another are unsubstituted or alkyl-substituted C_1 - C_5 alkylene.

Representative compounds falling with formula (IA) are propyl benzyl phthalate, butyl benzyl phthalate (Santicizer 160, Solutia), pentyl benzyl phthalate, hexyl benzyl phthalate, heptyl benzyl phthalate (Santicizer 261, Solutia), octyl benzyl phthalate, nonyl benzyl phthalate, decyl benzyl phthalate and C_1 - C_3 alkyl-substituted compounds thereof. Another commercially available alkyl benzyl phthalate is Santicizer 278, Solutia). Representative compounds falling with formula (IB) are dipentyl phthalate, dihexyl phthalate, diheptyl phthalate, dioctyl phthalate, dinonyl phthalate, didecyl phthalate and C_1 - C_3 alkyl-substituted compounds thereof. Another commercially available alkyl benzyl phthalate is Santicizer 278, Solutia.

Representative compounds falling with formula (IIA) are pentyl diphenyl phosphate, hexyl diphenyl phosphate, heptyl diphenyl phosphate, octyl diphenyl phosphate, nonyl diphenyl phosphate and C_1 - C_3 alkyl substituted compounds thereof, such as isodecyl diphenyl phosphate (Santicizer 148) and 2-ethylhexyl diphenyl phosphate (Santicizer 141, Solutia) and Santicizer 2148 (Solutia). Representative compounds falling with formula (IIB) are triphenyl phosphate and tricresyl phosphate.

Representative monomers falling with formula (III) are tris(methoxymethyl)phosphate, tris(ethoxymethyl)phosphate, tris(propoxymethyl)phosphate, tris(butoxymethyl)phosphate, tris(pentoxymethyl)phosphate; tris(2-methoxyethyl)phosphate, tris(2-ethoxyethyl)phosphate, tris(2-propoxyethyl)phosphate, tris(2-butoxyethyl)phosphate, tris(2-pentoxyethyl)phosphate; tris(methoxypropyl)phosphate, tris(ethoxypropyl)phosphate, tris(propoxypropyl)phosphate, tris(butoxypropyl)phosphate, tris(pentoxypropyl)phosphate; tris(methoxybutyl)phosphate, tris(ethoxybutyl)phosphate, tris(propoxybutyl)phosphate, tris(butoxybutyl)phosphate, tris(pentoxybutyl)phosphate; tris(methoxypentyl)phosphate, tris(ethoxypentyl)phosphate,

tris(propoxypentyl)phosphate, tris(butoxypentyl)phosphate and tris(pentoxypentyl)phosphate. Isodecyl diphenyl phosphate is the most preferred.

The casting composition of the present invention is a mixture of the isocyanate component and/or isocyanate functional prepolymer blend, an aromatic amine curative, selected plasticizer(s) and optionally additional polyether- and/or polyester-polyol, and/or diisocyanate dimer. The mixture preferably contains about 40-80% by weight of an isocyanate functional prepolymer with an isocyanate content between 4 and 15% formed by the reaction of toluene diisocyanate with at least one polyol; 7-20% by weight of an aromatic polyamine; 6-30% by weight of a phosphate or phthalate ester plasticizer having a vapor pressure at 25°C of less than 100 mPa, and optionally, 0-20% by weight of uretdione diisocyanate (poly-HDI). In a more preferred embodiments, the isocyanate functional prepolymer blend has a free toluene diisocyanate content below 0.4%, most preferably below 0.1%.

A wide variety of fillers, dyes, and pigments can be used in the formulations described above. Examples of suitable fillers are calcium carbonate, clays, talcs, zinc oxide, titanium dioxide, and silica. The amount of filler usually is in the range of 0 to 800 parts per hundred by weight, depending on the application for which the formulation is intended.

Stabilizers known in the art can also be incorporated into the composition. The stabilizers may be for protection during the life of the finished product against, for example, oxygen, ozone, and ultra-violet radiation. Stabilizers can also be incorporated to protect against thermooxidative degradation during processing. Antioxidants and UV inhibitors that would otherwise interfere with the urethane curing process should be avoided. Preferred antioxidants are sterically hindered phenolic compounds. Stabilizers such as organic phosphites are also useful. Preferred UV inhibitors are benzotriazole compounds.

For curing these prepolymers, the number of $-NH_2$ groups in the aromatic diamine component should be approximately equal to the number of $-NCO$ groups in the isocyanate component and/or prepolymer blend. A small variation is permissible but in general from about 80 to 110% of the stoichiometric equivalent should be used, preferably about 85 to 100%.

The castable compositions described herein can be prepared by mixing the selected polyol(s) chain extenders with the isocyanate component and/or isocyanate functional preopolymer. The resulting chain extended product can then be used immediately or stored under a nitrogen blanket until such time that a casting is desired. The selected aromatic amine curative can be separately combined with the plasticizer and stored. Once a casting is desired, the chain extended product is combined with the aromatic amine curative and plasticizer and poured into a mould for curing and post-curing, if necessary.

Applications for cast elastomers include, potting and encapsulants, pipe seals, athletic surfaces, binders, drop hammer tools, prototype parts, foundry core boxes, pattern plates, bumping tools, working models for the ceramic industry and other moulded articles resulting cast mouldings. The present invention, therefore, further relates to moulded articles resulting from the formulations described herein. Preferably, the moulded article is free of any fiber reinforcing materials, such as carbon fibers, and yet exhibits a Shore D hardness in excess of about 45, more preferably in excess of about 50.

The following examples are illustrative of the present invention and are therefore not intended as a limitation on the scope thereof.

Example 1

The following components are combined in a closed vessel at ambient pressure and temperature. The components are intimately mixed for a sufficient time to produce a homogenous mixture, at which time the physical properties are determined.

Components

ADIPRENE LF750D	76.86g
Isodecyl diphenyl phosphate	23.02g
BYK A530	0.12g (degassing aid)
Dimethylthiotoluene diamine	17.00g

At 25°C viscosity of mixture is 3500 - 4000 mPas

Properties after curing at 25°C for 7 days:

Shore D hardness (ISO 868) 50 - 53

Tensile strength (ISO527)	20 - 25 MPa
Elongation at break (ISO527)	150 - 200 %

Example 2

The components identified below are combined analogously to the methods described in example 1 to produce a homogenous mixture.

ADIPRENE LF750D	53.80g
DESMODUR N3400	23.06g
Isodecyl diphenyl phosphate	23.06g
BYK A530	0.08g (a degassing aid)
Dimethylthiolumene diamine	24g

At 25°C viscosity of mixture is 1000 mPas

Properties after curing at 25°C for 7 days:

Shore D hardness (ISO 868)	63 - 66
Tensile strength (ISO527)	25 - 30 MPa
Elongation at break (ISO527)	150 - 200 %

Example 3

Several plasticizers identified below were used in a mixture of 76.86 grams Adiprene LF750D, 0.1 grams BYK A530 and 17 grams dimethylthiolumene diamine. The mixtures are then introduced into a mould and cast into either a 400x75x100 mm bar or a 100 mm diameter circle having a 5 to 10 mm thickness. The mixtures are allowed to cure at ambient conditions for 16 hours. The resulting cured bars are demoulded and left on a flat surface. Distortion is shown when the ends of the cured bars curl upwards within three days of demoulding. The results of various plasticizers are shown below:

Plasticizer	Amount (g)	Evaporation rate	Vapor pressure at 25°C (mm Hg)	Vapor pressure at 25°C (mPa)	Distortion of cured specimens
gamma- butyrolacto ne	15.37		0.3	40000	Yes
Butyl glycol acetate	15.37		0.3	40000	Yes
Dibasic ester	12.3		0.08	11000	Yes
Propylene carbonate	15.37		0.08	11000	Yes
Dibutyl phthalate	23.06	44	0.0063	840	Slight
Ethylhexyl diphenyl phosphate	23.06	7.4	6×10^{-5}	8	No
Butyl benzyl phthalate	30.74	7.7	3×10^{-7}	4×10^{-2}	No
Isodecyl diphenyl phosphate	23.06	2.8	3×10^{-8}	4×10^{-3}	No
tributoxy ethyl phosphate	23.06	6.2	$10^{-4} - 10^{-6}$	0.13 - 13	No

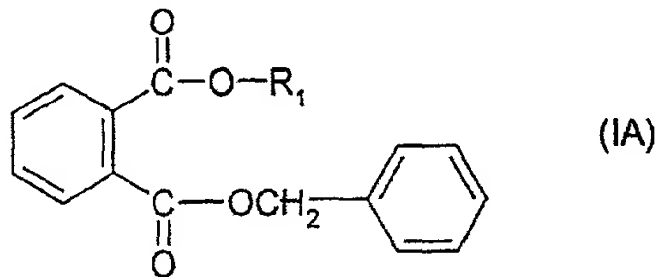
Preferred embodiments of the present invention relating to novel poly urethane/urea-forming compositions and methods for using the same have been described above. Those skilled in the art having the benefit of the teachings presented in the foregoing will recognize modifications and other embodiments. Therefore, it is understood that the

invention is not limited to the specific embodiments disclosed herein, and that modifications and other embodiments are intended to be within the scope of the appended claims.

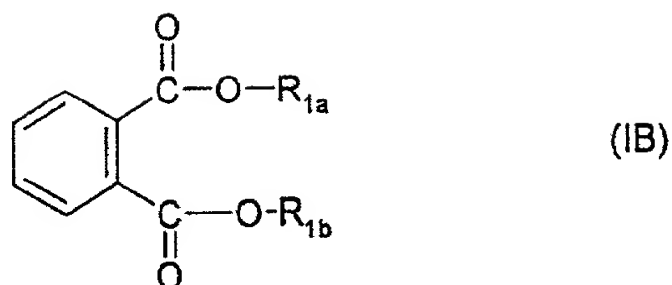
PCT/EP99/10040

Claims:

1. A poly urethane/urea-forming casting composition that can be cast and cured at temperatures between 15 and 35°C comprising a reaction mixture of (a) an isocyanate component or an isocyanate functional prepolymer having at least two isocyanate groups per molecule that contains or has been reacted with polytetramethylene glycol; (b) an aromatic amine curative; and (c) a phosphate ester or phthalate ester having a vapor pressure of less than 100 mPa at 25°C.
2. A casting composition according to claim 1 wherein component (a) is a low free toluene diisocyanate prepolymer blend having a free toluene diisocyanate content below 0.4%.
3. A casting composition according to claim 2 wherein the reaction mixture comprises a prepolymer blend that is a reaction mixture of an organic diisocyanate and polytetramethylene glycol.
4. A casting composition according to claim 1 wherein the aromatic amine curative is selected from the group consisting of diethyl toluene diamine, tertiary butyl toluene diamine, dimethylthiotoluene diamine, , and 1,2-bis(2-aminophenylthio)ethane.
5. A casting composition according to claim 4 wherein the aromatic amine curative is dimethylthiotoluene diamine.
6. A casting composition according to claim 5 further comprising a polyether- and/or polyester polyol having a number average molecular weight of at least 250.
7. A casting composition according to claim 1 wherein the phosphate ester or phthalate ester is represented by formulae (IA), (IB), (IIA), (IIB) or (III):

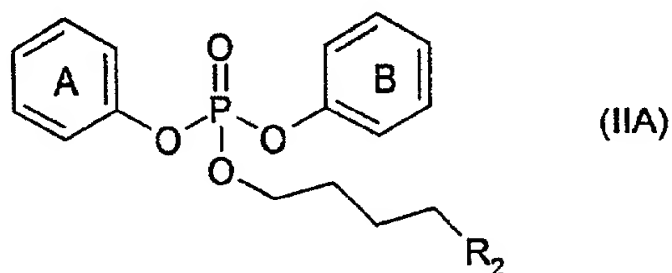


wherein R₁ is unsubstituted or alkyl-substituted C₃-C₁₂alkyl,

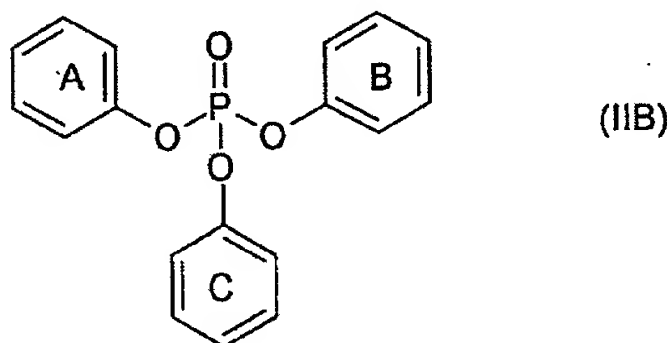


wherein R_{1a} and R_{1b} , independently of one another are unsubstituted or alkyl-substituted C_5 - C_{12} alkyl;

or phosphate esters according to formulae (IIA) and (IIB)

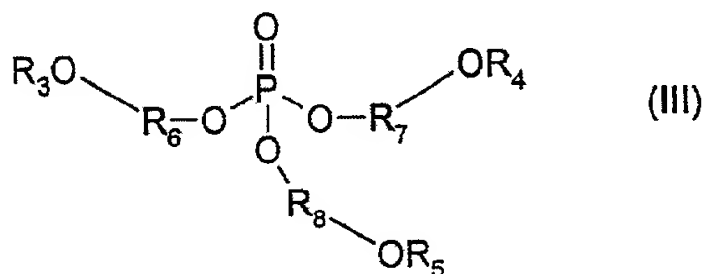


wherein R_2 is hydrogen or unsubstituted or alkyl-substituted C_2 - C_6 alkyl, and aromatic rings A and B independently of one another can have one or more alkyl substitutions;



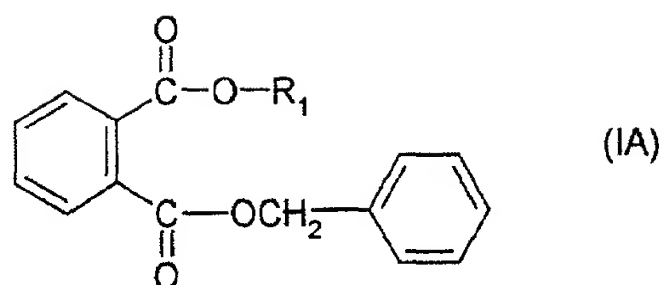
wherein aromatic rings A, B and C independently of one another can have one or more alkyl substitutions,

or formula (III)

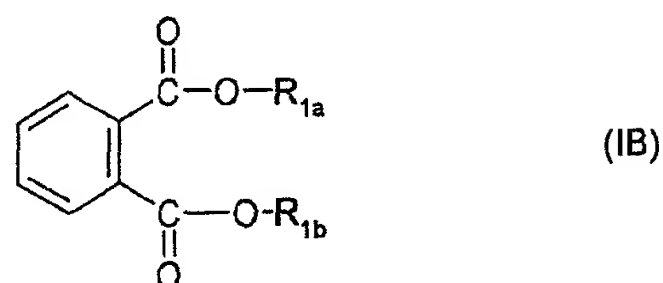


wherein R_3 , R_4 , R_5 , independently from one another, are unsubstituted or alkyl substituted C_1 - C_5 alkyl and R_6 , R_7 and R_8 are independently of one another are unsubstituted or alkyl-substituted C_1 - C_5 alkylene.

8. A casting composition according to claim 1 wherein component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.
9. A casting composition according to claim 5 wherein component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.
10. A cast polyurethane obtainable by a process comprising reacting (a) an isocyanate component or an isocyanate functional prepolymer having at least two isocyanate groups per molecule that contains or has been reacted with polytetramethylene glycol; (b) an aromatic amine curative; and (c) a phosphate ester or phthalate ester having a vapor pressure of less than 100 mPa at 25°C.
11. A polyurethane casting according to claim 10 wherein component (a) is a low free toluene diisocyanate prepolymer blend having a free toluene diisocyanate content below 0.4%.
12. A polyurethane casting according to claim 11 wherein component (a) is a prepolymer blend resulting from a reaction mixture of an organic diisocyanate and polytetramethylene glycol.
13. A polyurethane casting according to claim 10 wherein the aromatic amine curative is selected from the group consisting of diethyl toluene diamine, tertiary butyl toluene diamine, dimethylthiotoluene diamine, and 1,2-bis(2-aminophenylthio)ethane.
14. A polyurethane casting according to claim 13 wherein the aromatic amine curative is dimethylthiotoluene diamine.
15. A polyurethane casting according to claim 14 wherein the process further comprising combining a polyether- and/or polyester polyol having a number average molecular weight of at least 250 with components (a), (b) and (c).
16. A polyurethane casting according to claim 10 wherein the phosphate ester or phthalate ester is represented by formulae (IA), (IB), (IIA), (IIB) or (III):

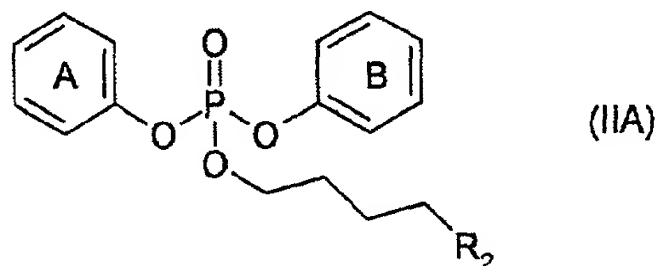


wherein R_1 is unsubstituted or alkyl-substituted C_3 - C_{12} alkyl,

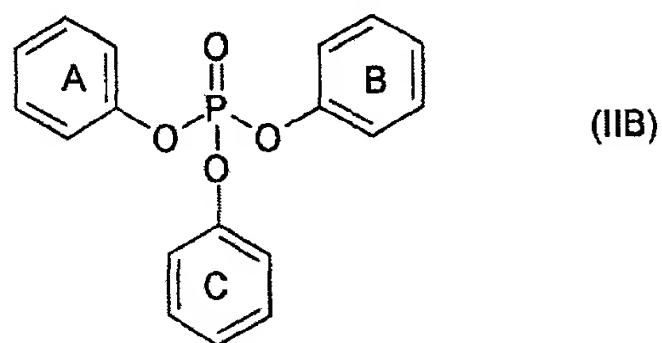


wherein R_{1a} and R_{1b} , independently of one another are unsubstituted or alkyl-substituted C_5 - C_{12} alkyl,

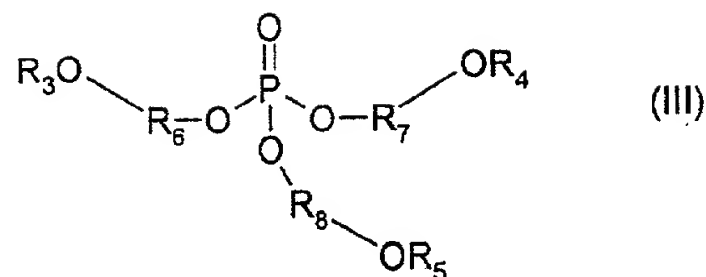
or phosphate esters according to formulae (IIA) and (IIB)



wherein R_2 is hydrogen or unsubstituted or alkyl-substituted C_2 - C_6 alkyl, and aromatic rings A and B independently of one another can have one or more alkyl substitutions;



wherein aromatic rings A, B and C independently of one another can have one or more alkyl substitutions,
or formula (III)



wherein R_3 , R_4 , R_5 , independently from one another, are unsubstituted or alkyl substituted $\text{C}_1\text{-C}_5$ alkyl and R_6 , R_7 and R_8 are independently of one another are unsubstituted or alkyl-substituted $\text{C}_1\text{-C}_5$ alkylene.

17. A polyurethane casting according to claim 10 wherein component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.

18. A polyurethane casting according to claim 14 wherein component (c) is tributoxyethyl phosphate or isodecyl diphenyl phosphate.

19. A process for curing a poly urethane/urea-forming composition according to claim 1 comprising contacting said poly urethane/urea-forming composition at a temperature between 15 and 35°C with (a) an aromatic amine curative having at least two primary amine groups; and (b) a plasticizer having a vapor pressure of less than 100 mPa at 25°C with said poly urethane/urea-forming composition.

**UTILITY DECLARATION
AND POWER OF ATTORNEY
Utility Application**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

THERMOSETTING POLY URETHANE/UREA-FORMING COMPOSITIONS

the specification of which

- ☐ is attached hereto OR
☐ was filed on _____ as United States Application Serial No. ____ OR
☒ was filed on 17/12/99 as PCT International Application No. PCT/EP 99/10040
☐ and was amended on ____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment(s) referred to above.

I acknowledge the duty to disclose information which is material to the patentability of this application in accordance with Title 37, Code of Federal Regulations, § 1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) or § 365(b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT international application which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or of any PCT international application having a filing date before that of the application on which priority is claimed.

Prior Foreign Application Number(s)	Country	Date of Filing (day/month/year)	Priority Claimed	
			Yes	No
60/112,656 /	USA /	17/12/98 /	<input checked="" type="checkbox"/>	<input type="checkbox"/>
			<input type="checkbox"/>	<input type="checkbox"/>

I hereby claim the benefit under Title 35, United States Code § 119(e) of any United States provisional application(s) listed below.

Application Number(s)	Filing Date (day/month/year)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s), or § 365(c) of any PCT international application designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

U.S. Parent Application Number	PCT Parent Number	Parent Filing Date (day/month/year)	Status-Patented, Pending or Abandoned

POWER OF ATTORNEY: As a named inventor, I hereby appoint as my attorneys and/or agents, with full power of substitution and revocation, to prosecute this application and transact all business in the United States Patent and Trademark Office, and in countries other than the United States, and to do all things necessary or appropriate therefor before any competent International Authorities in connection with any international patent application(s) corresponding to the above-identified invention application, all of the registered practitioners identified by Customer Number 22249:



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Please direct all inquiries to Kristin H. Neuman, Esq., at the above Customer Number. Ms. Neuman's direct line is (914) 681-8851 ext. 4603.

I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Title 18, United States Code, § 1001 and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

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